



## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

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<b>(21) International Application Number:</b> PCT/US97/13580 <b>(22) International Filing Date:</b> 31 July 1997 (31.07.97)  <b>(30) Priority Data:</b> 08/691,106 1 August 1996 (01.08.96) US  <b>(71) Applicant:</b> EXXON CHEMICAL PATENTS INC. [US/US]; 5200 Bayway Drive, Baytown, TX 77520-5200 (US).  <b>(72) Inventors:</b> MIDDLESWORTH, Jeffrey, A.; 116 S. Maple Street, Wauconda, IL 60084 (US). BRADY, Kevin, A.; 317 Claire Lane, Cary, IL 60013 (US).  <b>(74) Agents:</b> SHER, Jaimes et al.; Exxon Chemical Company, P.O. Box 2149, Baytown, TX 77522-2149 (US).		<b>(81) Designated States:</b> AU, CA, JP, European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).  <b>Published</b> <i>With international search report.</i> <i>With amended claims.</i>
<b>(54) Title:</b> BREATHABLE FILM/NWF-LAMINATES HAVING HIGH WVTR PREPARED FROM MELT EMBOSSED POLY-OLEFIN/FILLER PRECURSOR FILMS		
<b>(57) Abstract</b>  Films, made of polyethylenes and fillers, and articles made therefrom greater WVTR than previously available films based on conventional Zeigler-Natta based polyethylenes. The polyethylenes are produced in a metallocene-catalyzed production process. The films may be made by a cast film process, and may be made in widely varying filler content, generally polyethylene to filler ratios of 30/70 to 70/30. The metallocene based polyethylenes when combined with filler also permit the extrusion of thinner films leading to lighter weight and softer films. The m-polyethylenes utilized for making such films typically have a Composition Distribution Breadth Index above 50 %, an $M_w/M_n$ below 3, and an $M_z/M_w$ below 2. The films may be used advantageously in composite structures with fabrics (either woven or non-woven) to fabricate such articles as house-wrap.		

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**BREATHABLE FILM/NWF-LAMINATES HAVING HIGH WVTR PREPARED FROM MELT EM-  
BOSSSED POLYOLEFIN/FILLER PRECURSOR FILMS****TECHNICAL FIELD**

5           This invention relates generally to polyolefin films having greatly increased WVTR and methods of making same. More specifically this invention is directed toward filled polyethylene films having increased WVTR at a given filler loading, and a given set of process conditions.

**BACKGROUND**

10           Preparation of films having good WVTR from highly filled polymers, usually polyolefins, are known. In the past a combination of a polyolefin, usually a polyethylene, with a filler, usually  $\text{CaCO}_3$ , while very useful and widely used as a film with good WVTR, usually in combination with non-woven polymers (for use in diapers, adult incontinence devices, feminine hygiene articles, housewrap  
15           composites, roofing materials and the like), have had some limitations that were well known in the industry.

          Among these limitations are a practical limitation in thickness (also expressed as basis weight) in that conventional Ziegler-Natta catalyzed polymers, more specifically linear low density polyethylene (LLDPE) highly filled film  
20           formulations could not generally be drawn down below 3 mils. The most obvious problem with such a limitation is that the user of the film could not make a product utilizing a lower thickness film, meaning that the cost of the film (usually sold on a weight basis) might have been higher than the application necessitated. A less  
25           obvious issue is that at lower thicknesses, for the same density resin at the same filler loading, the product would be relatively softer than higher thicknesses, an attribute of importance in any article that comes in contact with humans, such as apparel.

          Another limitation of previous polyethylene/filler films is that for a given filler loading, with conventional Z-N catalyzed polyethylene resins, is WVTR,  
30           limited (on the upper end) by the amount of post-extrusion orientation that could be practically achieved. Additionally, the imperfections often found in conventional

Z-N resins and films, such as gels, made reaching and maintaining a high rate of production difficult, and a high level of orientation might often lead to breaks, holes, or tear offs in the film leading to lower prime production rates.

Yet another limitation of the conventional Z-N filled and oriented films is related to both WVTR and production rates. Specifically, with a given conventional filled polyethylene, to attain a certain WVTR, a certain filler loading had to be used. In general, within limits, the higher the filler loading, the more difficult to process ( the above referenced production problems such as large void creation and tear offs are exacerbated by a higher filler loading, as the film maker seeks to maximize production rates).

US 4,777,073 suggests a permeability and strength of polyethylene/filler combinations may be attained by combining a LLDPE described as being made using a Zeigler-Natta or chromium catalysts, with fillers such as  $\text{CaCO}_3$  present in the LLDPE from 15 to 35 percent by volume which is equivalent to 34-62% by weight.

There is a commercial need therefore for a polyethylene filler combination that will give a higher WVTR at a given filler loading, at an equivalent thickness. There is a similar need for a polyethylene filler combination that can deliver equivalent WVTR at lower filler loadings and can be made at a lower basis weight, than a conventional Z-N polyethylene/filler combination.

### SUMMARY

We have discovered that making a film from a polyethylene/filler combination using a metallocene catalyzed polyethylene, surprisingly and unexpectedly provides the ability to achieve a substantially higher WVTR (at comparable filler loading and thickness), a lower thickness (or basis weight) (at comparable filler loading and orientation), and can achieve an equivalent WVTR at lower filler loadings (improving processability) when compared to conventional Z-N polyethylene/filler combination.

The metallocene catalyzed polyethylenes (m-polyethylene) will have a molecular weight distribution (defined as the ratio of weight to the number average molecular weight  $M_w/M_n$ ) generally less than 3, preferably less than 2.5.

The drawdown of a filled m-polyethylene will be more than 10, preferably more than 20, more preferably more than 30 percent less than the ultimate drawdown of a filled Z-N polyethylene, where the relationship in the filled Z-N polyethylene between the filler amount and basis weight (minimum) for films follow the general equation:

$$W = 2.10 + 0.380 (\text{weight \% CaCO}_3)$$

where W is the minimum basis weight in g/m<sup>2</sup> in the film.

The relationship is at constant draw (orientation transverse direction or TD) of 2.7:1, line speed 340 feet per minute (fpm). For m-polyethylene filled formulations the following general equation applies:

$$W = 3.07 + 0.207 (\text{weight \% CaCO}_3)$$

Additionally the water vapor transmission rate (WVTR) of a filled m-polyethylene is at least 10 percent greater, preferably at least 20 percent, more preferably at least 30 percent greater than a filled Z-N polyethylene, at the same filler loading and thickness (basis weight), where the Z-N polyethylene/filler WVTR is described by the equation:

$$\text{WVTR} = -10,900 + 320 (\text{weight \% CaCO}_3)$$

where the WVTR is in g/m<sup>2</sup>/24 hours, measured at 37.8° C, 90% RH. While a film including a m-polyethylene and filler follows the general equation:

$$\text{WVTR} = -9967 + 358 (\text{weight \% CaCO}_3)$$

The relationship is at constant draw (orientation TD) of 2.7:1, line speed 340 feet per minute (fpm).

#### **BRIEF DESCRIPTION OF THE DRAWINGS**

The foregoing aspects, features and advantages of the present invention will become clearer and more fully understood when the following detailed description, and appended claims are read in conjunction with the accompanying drawings, in which:

Figure 1 illustrates the drawdown advantage of filled m-polyethylene over Z-N polyethylene with a plot of minimum basis weight in g/m<sup>2</sup> versus filler loading.

Figure 2 illustrates the WVTR advantage of m-polyethylene versus Z-N polyethylene in a plot of WVTR versus percentage of filler  $\text{CaCO}_3$  both at 2.7:1 draw ratio and 22 g/m<sup>2</sup> basis weight.

## **DETAILED DESCRIPTION**

### **Introduction**

This invention concerns certain polyethylene/filler films that will have high WVTR and the ability to be drawn down to low basis weights and methods for making same. Particularly useful in these films and methods will be m-polyethylenes.

In certain embodiments of the present invention films of m-polyethylene and filler can be made with lower amounts of filler and still attain substantially the same WVTR as previously known and used Z-N polyethylene/filler combinations (at higher filler loadings) are also contemplated. This invention further includes certain m-polyethylenes, their conversion into fabricated articles such as films, articles made from such films, and applications in which such articles having high WVTR combined with good physical properties are desirable. The resulting films, and film composites, (including coextruded and laminated films) have combinations of properties rendering them superior and unique to films or film composites previously available. The filled m-polyethylene films disclosed herein are particularly well suited for use in producing certain classes of high WVTR films, consumer and industrial articles using the films in combination with for instance, polymeric woven or non-woven materials. Such consumer articles include, but are not limited to diapers, adult incontinence devices, feminine hygiene articles, medical and surgical gowns, medical drapes, industrial apparel, building products such as "house-wrap", roofing components, and the like made using one or more of the films disclosed herein. Additionally the films having increased WVTR of the present invention may also be used in metallized films with a high WVTR, according to the disclosure of U.S. Patent 5,055,338, fully incorporated herein for purposes of U.S. Patent practice.

### **Housewrap**

Fabrics suitably laminated to the breathable film in the housewrap of the present invention include any high strength fabric which can be bonded to the breathable film without adversely affecting the water vapor permeability or the resistance to air permeability of the breathable film, i.e. the fabric must generally have a suitably open mesh to avoid substantially blocking the micropores of the breathable film. The fabric may be woven of any suitable material, but is preferably non woven polyolefin such as, for example, low density polyethylene, polypropylene, and preferably linear, low density polyethylene or high density polyethylene. The fabric should have an elongation (ASTM D1682) less than 30%: an Emendorf tear strength (ASTM D689) of at least 300 g, preferably at least 600 g and especially at least 900 g; and a break load (ASTM D1682) of at least 15 lb/in., preferably at least 25 lb/in., and especially at least 30 lb/in/. These fabrics are believed to be prepared from HDPE films having outer layers of ethylene vinyl acetate coextruded on either side of the HDPE or heat seal layers. The films are fibrillated, and the resulting fibers are spread in at least two transverse directions at a strand count of 6010 per inch. The spread fibers are then cross laminated by heat to produce a nonwoven fabric of 3-5 mils with equal MD and TD strength. These fabrics have excellent strength properties in both MD and TD for reinforcing the breathable film, an open structure to avoid substantially blocking the micropores of the breathable film when laminated thereto, and an outer layer of ethylene vinyl acetate copolymer for heat sealability.

The fabric and the breathable film are laminated together to form the breathable composite of the invention. The lamination may be effected by facing the film and the fabric together and applying heat and pressure. The laminating temperatures to which the film and fabric are exposed should be sufficient to achieve lamination, but should not be too high in order to avoid the flow of the film polymer into the microporous spaces and consequent reduction in water vapor transmissibility. In a preferred embodiment, the fabric is heated on a hot roller, preferably at 200° - 240° F., and then pressed, preferably at a pressure of 50 - 100 psi, into contact with the unheated film to bond the fabric and film into laminate.

Preferred fabrics are commercially available under the trade designation DD1001, CC-2001 and CC-3001 CLAF nonwoven HDPE Fabrics.

In an embodiment of our invention, the filled m-polyethylene films, when oriented after film formation, would surprisingly and unexpectedly have high WVTR when compared to a filled polyethylene film made using previously available Z-N catalyzed polyethylenes. Following is a detailed description of certain preferred m-polyethylenes, films, or film composites made using these m-polyethylenes and articles made from the films or film composites, that are within the scope of the present invention. Those skilled in the art will appreciate that numerous modifications to these preferred embodiments can be made without departing from the scope of the invention. For example, although films based on low density m-polyethylenes filled with  $\text{CaCO}_3$  are exemplified herein, the films may be made using combinations of m-polyethylenes with other polyolefins and with other fillers or filler combinations. To the extent my description is specific, it is solely for the purpose of illustrating preferred embodiments of my invention and should not be taken as limiting the present invention to these specific embodiments.

#### **Production of the Films**

Films contemplated by certain embodiments of the present invention may be made utilizing m-polyethylenes, by processes including, blown and cast, preferred is a cast film process. In such extrusion processes, the films of the present invention can be formed into a single layer film, or may be one layer or more of a multi-layer film or film composite. Alternatively, the m-polyethylene films described in this disclosure can be formed or utilized in the form of a resin blend where the blend components can function to modify WVTR, physical properties, draw-down sealing, cost, or other functions. Both blend components and functions provided thereby will be known to those of ordinary skill in the art. Films of the present invention may also be included in laminated structures. As long as a film, multi layer film, or laminated structure includes one or more m-polyethylene/filler film layers having the WVTR, or draw-down, and the like of the film, and the  $M_w/M_n$ , CDBI and the like of the m-polyethylene, in the ranges described herein, it will be understood to be contemplated as an embodiment of the present invention.

Polyolefin Component

The polyolefin component can be any film forming polyolefin or polyolefin blend, as long as the majority of the polyolefin component is a polyolefin with the following features:

5		preferred	more preferred	most preferred
	Mw/Mn	< 3	<2.5	
	CDBI	>50%	>60%	>65%
	Mz/Mn	<2		

Generally these ranges dictate the use of a metallocene catalyzed polyolefin, preferred is a m-polyethylene, preferably a linear low density m-polyethylene with a density in the range of from 0.90-0.940, preferred 0.910-0.935, more preferred 0.912-0.925 g/cc. Densities referred to herein will generally be polymer or resin densities, unless otherwise specified.

There is a wide variety of commercial and experimental m-polyethylene resins useful in the manufacture of films included in certain embodiments of the present invention. A non-inclusive list is found below along with the general bulk resin properties as published:

**TABLE A**

20	Commercial Designation	Density (g/cm <sup>3</sup> )	Melt Index/ Melt Flow (g/10 min.)	Type
25	Exceed® 103 (now 350L65 or 350D60)*	0.917	1.0	eth/hexene
	Exceed® 301 now 357C80*	0.918	3.4	eth/hexene
	Exceed® 377D60*	0.922	1.0	eth/hexene
	Exceed® 109*	0.925	0.75	eth/hexene
	Exact® 3028*	0.900	1.2	eth/butene
30	Exceed® 357C32*	0.917 g/cc	3.4	
	Exceed® 363C32	0.917 g/cc	2.5	
	ECD-401	0.917 g/cc	4.5	
	Exceed® 377D60	0.922 g/cc	1.0	
35	Exceed® 399L60	0.925 g/cc	.75	

\*available from Exxon Chemical Co. Houston, TX, USA.

\*The Exceed® 357C32 is the same resin grade as the ECD-112 and ECD-115 used in the experiments.

It will be understood that in general we contemplate that a large number of m-polyethylenes will be useful in the techniques and applications described herein. Included components: ethylene-1-butene copolymers, ethylene-1-hexene copolymers, ethylene-1-octene copolymers, ethylene-4-methyl-1-pentene copolymers, ethylene dodecene copolymers, ethylene-1-pentene copolymers, as well as ethylene copolymers of one or more C4 to C20 containing alpha-olefins, diolefins, and combinations thereof. A nonexclusive list of such polymers; ethylene, 1-butene, 1-pentene; ethylene, 1-butene, 1-hexene; ethylene, 1-butene, 1-octene; ethylene, 1-butene, decene; ethylene, 1-pentene, 1-hexene; ethylene, 1-pentene, 1-octene; ethylene, 1-pentene, decene; ethylene, 1-octene; 1-pentene; ethylene 1-octene, decene; ethylene, 4-methyl-1-pentene, 1-butene; ethylene 4-methyl-1-pentene, 1-pentene; ethylene, 4-methyl-1-pentene, 1-hexene; ethylene 4-methyl-1-pentene, 1-octene; ethylene, 4-methyl-1-pentene, decene. Included in the ethylene copolymers will be one or more of the above monomers included at a total level of 0.2 to 6 mole percent, preferably 0.5 to 4 mole percent, or such mole percents consistent with the resin densities contemplated.

Definitions and methods of determination of CDBI may be found in U.S. 5,008,204 which is fully incorporated by reference herein for purposes of U.S. patent practice.

The resin and product properties recited in this specification were determined in accordance with the following test procedures. Where any of these properties is referenced in the appended claims, it is to be measured in accordance with the specified test procedure.

TABLE B

Property	Units	Procedure
Melt Index	dg/min	ASTM D-1238(E)
Density	g/cc	ASTM D-1505
WVTR	g/m <sup>2</sup> /day	described herein
Gurley	seconds	described herein

#### FILLER

Fillers useful in this invention may be any inorganic or organic material having a low affinity for and a significantly lower elasticity than the polyolefin

component. Preferably the filler should be a rigid material having a non-smooth hydrophobic surface, or a material which is treated to render its surface hydrophobic. The preferred mean average particle size of the filler is between 0.5-5 microns for films generally having a thickness of between 1-6 mils prior to stretching. Examples of the inorganic fillers include calcium carbonate, talc, clay, kaolin, silica, diatomaceous earth, magnesium carbonate, barium carbonate, magnesium sulfate, barium sulfate, calcium sulfate, aluminum hydroxide, zinc oxide, magnesium hydroxide, calcium oxide, magnesium oxide, titanium oxide, alumina, mica, glass powder, zeolite, silica clay, etc. Calcium carbonate is particularly preferred for low cost, whiteness, inertness, and availability. The inorganic filler such as calcium carbonate are preferably surface treated to be hydrophobic so that the filler can repel water to reduce agglomeration of the filler. Also, the surface coating should improve binding of the filler to the polymer while allowing the filler to be pulled away from the polyolefin under stress. A preferred coating is calcium stearate which is FDA compliant and readily available. Organic fillers such as wood powder, and other cellulose type powders may be used. Polymer powders such as Teflon ® powder and Kevlar ® powder can also be used.

The amount of filler added to the polyethylene depends on the desired properties of the film including tear strength, water vapor transmission rate, and stretchability. However, it is believed that a film with good WVTR generally cannot be produced as is taught herein with an amount of filler less than 20 percent by weight of the polyolefin/filler composition.

The minimum amount of filler is needed to insure the interconnection within the film of voids created at the situs of the filler particularly by the stretching operation to be subsequently performed on the precursor film. Further, it is believed that useful films could not be made with an amount of the filler excess of 70 percent by weight of the polyolefin/filler composition. Higher amounts of filler may cause difficulty in compounding and significant losses in strength of the final breathable film.

While a broad range of fillers has been described at a broad range of inclusion parameters based on weight percentages, other embodiments are

contemplated. For instance, fillers with much higher or much lower specific gravities may be included in the polyolefin at amounts outside the weight ranges disclosed, they will be understood to be contemplated as embodiments of our invention as long as the final film, after orientation has WVTR or drawn down similar to that described herein.

#### STRETCHING OR ORIENTING AND HEAT SETTING

Final preparation of a breathable film is achieved by stretching the filled m-polyethylene precursor film to form interconnected voids. Stretching or "Orientation" of the film may be carried out monoaxially in the machine direction (MD) or the transverse direction (TD) or in both directions (biaxially) either simultaneously or sequentially using conventional equipment and processes following cooling of the precursor film.

Film orientation may also be carried out in a tentering device with or without MD orientation to impart TD orientation to the film. The film is gripped by the edges for processing through the tentering device.

Stretching of melt embossed precursor films with a tentering device at a film speed of 200-500 per minute produces breathable films having the desired water vapor permeability. The resulting films had a greater permeability in the areas of reduced thickness in comparison to the areas of greater thickness.

A range of stretching ratios from 2:1 to 5:1 prove satisfactory for MD stretching with a ratio of 4:1 being preferred. A range of stretching ratios of 2:1 to 5:1 prove satisfactory for TD stretching with a ratio of 3:1 being preferred.

It is preferred that tension be maintained on the film during the heat setting and cooling to minimize shrinkback. Upon cooling to ambient temperature (i.e., room temperature) or near ambient, the holding force may be released. The film may contract somewhat (snapback) in the TD but will retain a substantial portion of its stretched dimension.

Heat setting can be accomplished by maintaining the film under tension in the stretched condition at the heat setting temperature for 1-2 minutes. Preferably, however, the heat setting and cooling is carried out while permitting the film to

contract slightly, but still under stress. The controlled shrinkback of from 5 to 30%, preferably between 15 and 25%, of the maximum stretched width has given particularly good results in eliminating storage shrinkage.

**Properties of films produced from the resins**

5 **WVTR**

In an embodiment of the present invention, certain films and articles made therefrom have higher WVTR than previously thought possible. The WVTR of such films should be above 100 g/m<sup>2</sup>/day @ 37.8° C, 90% RH, preferably above 1000, more preferably above 3000 g/m<sup>2</sup>/day @ 25° C. This can be seen in Figure 2  
10 which illustrates the WVTR advantage of m-polyethylene versus Z-N polyethylene in a plot of WVTR versus percentage percentage of filler CaCO<sub>3</sub>.

In general the films of embodiments of the present invention will have a much higher WVTR at the same filler loading than previously known Z-N polyethylene based filled films. Specifically, the inventive films will have a WVTR  
15 at least 10% higher than the WVTR of the comparative films described by the equation: 
$$\text{WVTR} = -10,900 + 320 (\text{weight \% CaCO}_3)$$

In another embodiment of our invention a m-polyethylene/filler combination film can be stretched (oriented or tentered in the TD) less than a Z-N polyethylene combination film, and still achieve substantially the same WVTR (at generally the  
20 same filler loadings). This is a significant advantage to a film maker because the greater the orientation, the greater the chance for a film imperfection to be magnified, potentially causing a catastrophic failure (break).

It is not beyond the scope of embodiments of my invention to blend the m-polyolefins to form the films of the invention with other materials such as other  
25 linear polyethylenes (HDPE, MDPE, LLDPE), low density polyethylene (LDPE), polypropylene (PP) (homopolymers and copolymers), polybutene-1 (PB), ethylene vinyl acetate (EVA), or other ethylene polar comonomer copolymers and the like to fabricate useful articles. Such potential blend polyolefins may be conventional Zeigler-Natta catalyzed, chromium catalyzed, free radical initiated, and the like.  
30 However, the WVTR of the layer or layers intended to impart WVTR should generally be within limits disclosed above. Additionally, any blend component or

components additive or additives should be chosen such that the desired WVTR of the film remains at or above the targeted or desired value. Any blend should preferably contain a majority of m-polyethylene as the polyolefin component, specifically greater than 50 weight percent, preferably greater than 60 weight percent, more preferably greater than 70 (75?) percent, based on the total weight of the polyolefin

#### Definitions and Test Protocols

Value	Units	Definition or Test
Density	g/cm <sup>3</sup>	ASTM D-792
CDBI	%	*Definitions test determination contained in this application
Molecular weight distribution	none	

#### TEST METHODS

##### 10 Water Vapor Transmission Rate

The WVTR test measures the quantity of water vapor that is able to pass through a film. A Mocon Permatran W-1 unit is used to measure WVTR by passing a stream of dry air across the surfaces of the film. The dry air picks up moisture that has passed, from wet pads underneath the film, through to the top surface.

The moisture level is measured by an infrared (IR) detector and converted to a voltage which can be measured on a chart recorder. The procedure also includes:

- a) Punching out a die cut hole in an aluminum foil mask,
- 20 b) Cutting off two opposing corners of the mask,
- c) Peeling paper backing off of mask,
- d) Cutting 2" x 2" squares of film and place them over the hole in the mask,
- e) Putting the paper backing back on the foil mask, then
- 25 f) Placing the masked sample in the test cell with the aluminum side up.

The chart recorder reading is multiplied by 100 to give the WVTR value.

Gurley Porosity

Teledyne Gurley Model 4190 Porosity Tester with sensitivity attachment is used. With the procedure as follows:

- a) Cutting a strip of film (~2" wide) across the entire web width,
- 5 b) Inserting a film sample to be tested between orifice plates,
- c) Setting the sensitivity adjustment on "5",
- d) Turning the inner cylinder so that the timer eye is vertically centered below the 10cc silver step on the cylinder,
- e) Resetting the timer to zero,
- 10 f) Pulling the spring clear of the top flange and releasing the cylinder,

When the timer stops counting, the test is completed. The number of counts is multiplied by 10 and the resulting number is "Gurley seconds per 100cc".

It will be appreciated by those of ordinary skill in the art that the films of m-polyethylene resins of certain embodiments of the present invention, can be  
15 combined with other materials, depending on the intended function of the resulting film.

Other methods of improving and/or controlling WVTR properties of the film or container may be used in addition to the methods described herein without departing from the intended scope of my invention. For example, mechanical  
20 treatment such as micro pores.

DRAWDOWN

Embodiments of the present invention offer a significant and unexpected improvement in the ability for the formulations to be drawn down. Specifically, using conventional Z-N polyethylenes, a lower limit of 2.5, more practically 3.5  
25 mils has routinely been observed (as extruded) upstream, i.e. before orientation. By contrast, films of embodiments of the present invention, may be drawn down to a practical limit of 2 mils, providing a significant advantage in terms of either economics or a combination of economics and softness. The softness comes about due to the decreased modulus of the lower thickness. Ultimate drawdown  
30 is defined as minimum gage (or basis weight) before the onset of draw resonance at a given extruder rate (e.g., lb/hr).

The films of embodiments of the present invention will have ultimate drawdown more than 20%, preferably 25%, more preferably 30% less than that of filled Z-N polyethylene which, from Figure 2 has an ultimate drawdown described by the general formula:

$$W = 2.1 + 0.380 (\text{weight \% CaCO}_3)$$

5     **EXAMPLES**

All polyethylene/filler materials were stabilized to diminish the effects of extrusion.

Orientation of all the following examples was performed at a 2.7:1 draw ratio, at 35° fpm, 150-220° F tenter temperature, 180-230° F annealing temperature.

10     **Example 1 - 3**

Examples 1 - 3 were fabricated from Escorene™ LL 3003.09 on a 6 inch Marshall & Williams cast extrusion line at normal processing conditions processing conditions listed in Table 1a.

15     Example 1 used a 50/50 weight ratio of the polyethylene to CaCO<sub>3</sub>, while examples 2-3 used a 65/35 ratio of polyethylene to filler all films were subsequently oriented (TD) to three different basis weights as seen in Table 1.

**Examples 4-9**

20     Examples 4-9 were fabricated from Exceed™ ECD-112, under the same processing conditions as examples 1-3. Examples 4-6 used a 50/50 weight ratio of the polyethylene to CaCO<sub>3</sub>, while examples 7-9 used a 65/35 ratio of polyethylene to filler. All films were subsequently oriented (TD) to three different basis weights as seen in Table 2.

25     From the data in Table 1 for each of these examples run, it can be seen that in Example 1 and 2; as filler level goes down, WVTR goes down dramatically, and as seen from example 3 a lower basis weight only marginally increases the WVTR of the film with a higher percentage of polyethylene. By contrast, from table 2 for examples 4-9, a much higher WVTR is achieved by the same filler loading and basis weight, than for the films of examples 1-3, moreover, while a higher percentage of polyethylene in the formulation (examples 4-6 vs. 7-9) generates a

30

diminution of WVTR, the percentage is far lower than that experienced for the Z-N polyethylene of examples 1-3 (95% reduction vs. 68% reduction)

#### Examples 10 - 15

5 Examples 10-15 are run as in Example 4-9, but the polyolefin component was a blend of LD-202 (12-MI, 0.917 g/cc low density polyethylene available from Exxon Chemical Co.) and ECD112. As can be seen from the data in Table 3, at the same basis weight Examples 4-6, and 7-9, the corresponding films of Exmples 10-15 had somewhat lower, but still acceptable WVTR. Also of note is Example 15 which was the lowest basis weight attainable in this series (1-15) of examples 10 (again orientation was TD).

#### Examples 16-23

Examples 16-23 were extruded similar conditions to the previous examples, into two (2) thickness of precursor (before orientation) film (4.5 and 6 mils) and oriented in the MD at 175° F. While WVTR results for this set of examples appear 15 to be substantially the same for both metallocene and Z-N polyethylenes, it is anticipated that when the orientation speed is increased, the m-LLDPE will show improved WVTR, over the Z-N-LLDPE, just as found in the TD orientation in examples 1-15. The results are shown in Tables 4 and 5.

#### Examples 24-25

20 Examples 24 and 25 were extruded under substantially the same conditions as the previous examples. Examples 24 is substantially the same in polyethylene/filler content as example 4 and example 24 is substantially the same make-up as example 1.

Example 24 was drawn (oriented) at a 2.7:1 draw ratio, while example 25 25 was drawn at a 3.8:1 ratio. These examples show that the m-LLDPE at a lower (28%) draw ratio than the Z-N LLDPE, example 24 has generally the same WVTR. The results are shown in Table 6.

While the present invention has been described and illustrated by reference 30 to particular embodiments thereof, it will be appreciated by those of ordinary skill in the art that the invention lends itself to variations not necessarily illustrated herein. For example, it is not beyond the scope of this invention to include

additives with the claimed films or to blend resins to form the claimed films with other polymers or laminate the claimed films to other materials such as polymer non-wovens and the like. For this reason, then, reference should be made solely to the appended claims for purposes of determining the true scope of the present invention.

**TABLE 1**  
**ORIENTED FILM PROPERTIES**  
**LL 3003.09 Based Samples**

PROPERTIES	Example 1	Example 2	Example 3
Basis Wt., g/m <sup>2</sup>	22.1	22.5	18.7
Yield, yd <sup>2</sup> /lb.	24.6	24.1	29.0
Emb. Cal., mils	1.17	1.13	.98
Gurley, seconds	1137	Off-Scale	Off-Scale
WVTR, g/m <sup>2</sup> /24	5100	300	500
MD Tear, g	473	486	386
TD 170° F	9	8.5	7.8
Opacity, %	59.5	39.1	38.1
MD 10%, g/in	319.8	417.9	392.0
MD 25%, g/in	352.1	429.6	414.1
MD Ult., g/in	456.2	494.4	492.3
MD Elg., %	343.8	340.8	358.6
TD 10%, g/in	688.0	900.4	728.0
TD 25%, g/in	1092	1391	1134
TD Ult., g/in	1725	2025	1842
TD Elg., %	127.1	131.6	136.5
DR Limit g/m <sup>2</sup>	21.1	15.4	--

- The "DR Limit" also known as "Ultimate Drawdown" is the basis weight at which we first observed draw resonance. The DR probe was conducted with the fpm fixed at 340 and the extruder RPM reduced gradually until the onset of draw resonance.

**TABLE 1a**  
**Extrusion Conditions**  
**(for 22 g/m<sup>2</sup> samples)**

CONDITION	Example 1	Example 2	Example 4	Example 7	Example 10	Example 14
Ext. RPM	23.0	26.4	19.1	21.6	29.5	31
Upstream, psi	3480	-	3740	3730	4630	4430
Down, psi	1310	-	1270	1160	1630	1490
Melt Temp., F	427	436	430	431	458	451
Up Width, "	23.4	23.3	21.8	21.8	31.1	30.4
Down Width, "	-	60.9	62.6	62.1	86.4	-
Cast Roll, fpm	318	341	340	340	339	340

draw ratio for all examples targeted at 2.7-1, (ratio of outlet width divided by inlet width)

**TABLE 2**  
**ORIENTED FILM PROPERTIES**  
**For Exceed™ ECD-112 Based Samples**

PROPERTIES	Example 4 22 g/m <sup>2</sup>	Example 5 18.5 g/m <sup>2</sup>	Example 6 15 g/m <sup>2</sup>	Example 7 22 g/m <sup>2</sup>	Example 8 18.5 g/m <sup>2</sup>	Example 9 15 g/m <sup>2</sup>
Basis wt g/m <sup>2</sup> (Target)						
Basis Wt., g/m <sup>2</sup>	22.7	18.6	15.2	22.8	19.2	14.8
Yield, yd <sup>2</sup> /lb.	23.9	29.2	35.7	23.8	28.3	36.7
Emb. Cal., mils	1.23	.96	.81	1.24	1.03	.77
Gurley, seconds	216	159	127	3608	2140	1095
WVTR, g/m <sup>2</sup> /24	7950	8350	8450	2575	3675	4010
MD Tear, g	400	360	330	418	405	292
TD 170° F	8.0	7.2	7.2	7.2	7.0	6.5
Opacity, %	66.2	62.3	59.1	51.6	48.3	44.9
MD 10%, g/in	299.6	221.6	191.9	434.4	369.6	288.1
MD 25%, g/in	383.3	247.1	213.0	435.0	368.2	285.3
MD Ult., g/in	496.9	323.6	296.5	501.6	411.9	304.7
MD Elg., %	327.5	290.0	331.2	293.1	276.4	271.4
TD 10%, g/in	737.3	623.6	513.7	932.9	836.4	678.6
TD 25%, g/in	1182	1003	851.8	1503	1342	1111
TD Ult., g/in	2261	1863	1574	2942	2689	2197
TD Elg., %	110.2	100.7	95.5	103.5	103.3	97.1
DR Limit g/m <sup>2</sup>	13.4	--	--	10.3	--	--

- The "DR Limit" is the basis weight at which we first observed draw resonance. The DR probe was conducted with the fpm fixed at 340 and the extruder RPM reduced gradually until the onset of draw resonance.

**TABLE 3**  
**ORIENTED FILM PROPERTIES**

For samples based on Exceed™ ECD-112 blended with LDPE (LD-202)

PROPERTIES	Example 10 37.5% ECD 12.5% LD 50% Calc 22 g/m <sup>2</sup>	Example 11 37.5% ECD 12.5% LD 50% Calc 18.5 g/m <sup>2</sup>	Example 12 37.5% ECD 12.5% LD 50% Calc 15 g/m <sup>2</sup>	Example 13 56.3% ECD 8.7% LD 35% Calc 22 g/m <sup>2</sup>	Example 14 56.3% ECD 8.7% LD 35% Calc 15.0 g/m <sup>2</sup>	Example 15 56.3% ECD 8.7% LD 35% Calc 12 g/m <sup>2</sup>
Basis Wt., g/m <sup>2</sup>	22.1	17.9	14.7	22.9	13.9	12.1
Yield, yd <sup>2</sup> /lb.	24.6	30.3	36.9	23.7	39.0	44.8
Emb. Cal., mils	1.08	.99	.73	1.11	.70	.62
Gurley, seconds	1345	814	398	13,703	6930	3717
WVTR, g/m <sup>2</sup> /24	4800	5725	5925	950	1100	2350
MD Tear, g	98	90	85	371	189	187
TD 170° F	6.0	6.8	7	6	7	7
Opacity, %	59.7	55.6	51.2	50.6	40.3	37.7
MD 10%, g/in	361.3	304.6	255.4	472.8	331.2	277
MD 25%, g/in	391.6	331.9	281.9	526.7	327.2	280.2
MD Ult., g/in	441.1	367.7	311.4	526.7	352.8	296.5
MD Elg., %	163.2	137.3	103.2	259.8	202.8	177.2
TD 10%, g/in	641.4	520.5	435.5	828.2	560	460.5
TD 25%, g/in	985	806.4	678.2	1294	888.4	733.4
TD Ult., g/in	1578	1307	1197	2569	1912	1408
TD Elg., %	97.8	96.6	104.2	110.0	113.2	103.3
DR Limit g/m <sup>2</sup>	<11.5	--	--	<6.4	--	--

- The "DR Limit" is the basis weight at which we first observed draw resonance. The DR probe was conducted with the fpm fixed at 340 and the extruder RPM reduced gradually until the onset of draw resonance.

**TABLE 4**  
**175° F Orientation**  
**4.5 mil precursor film**

PROPERTY	Example 16 50% CaCO <sub>3</sub> in ECD-115 4:1 Draw Ratio	Example 17 50% CaCO <sub>3</sub> in ECD-115 6:1 Draw Ratio	Example 18 50% CaCO <sub>3</sub> in LL3003.09 4:1 Draw Ratio	Example 19 50% CaCO <sub>3</sub> in LL3003.09 6:1 Draw Ratio
Basis Weight, g/m <sup>2</sup>	54.7	34.5	54.84	34.87
Embossed Caliper, mils	2.43	1.93	3.29	2.79
WVTR, g/m <sup>2</sup> /24 hours	6100	7950	6500	7250
Gurley Porosity, sec / 100 cc	855	307	581	379
MD Tensile at 5%, g/in	1094	1289	1084	1344
MD Tensile at 10%, g/in	2290	3034	2192	3041
MD Tensile at 25%, g/in	4540	--	3774	--
MD Tensile at Break, g/in	7273	7725	5085	6135
MD Elong. at Break, %	73.48	19.65	78.74	20.78
TD Tensile at 5%, g/in	201.1	102.4	178.7	104.9
TD Tensile at 10%, g/in	333.4	196.5	293.4	184.7
TD Tensile at 25%, g/in	432.9	317.6	375.9	263.9
TD Tensile at Break, g/in	568.6	318.1	482.8	276.9
TD Elong. at Break, %	350.1	241.7	315.7	228.5
MD Elmendorf Tear, grams	4	0	2	13.2
MD Shrink at 170° F, %	13.5	17.6	10.5	16.0
TD Shrink at 170° F, %	-3.0	-3.1	-3.8	-2.9

5 Note: All samples oriented with a 15 fpm inlet speed, 190° F annealing and 5% relaxation.

**TABLE 5**  
**175° F Orientation**  
**6.0 mil precursor film**

PROPERTY	Example 20 50% CaCO <sub>3</sub> in ECD-115 4:1 Draw Ratio	Example 21 50% CaCO <sub>3</sub> in ECD-115 6:1 Draw Ratio	Example 22 50% CaCO <sub>3</sub> in LL3003.09 4:1 Draw Ratio	Example 23 50% CaCO <sub>3</sub> in LL3003.09 6:1 Draw Ratio
Basis Weight, g/m <sup>2</sup>	63.19	47.95	65.72	44.47
Embossed Caliper, mils	3.30	2.68	3.20	2.55
WVTR, g/m <sup>2</sup> /24 hours	5450	7500	6250	7800
Gurley Porosity, sec / 100 cc	1151	363	541	282
MD Tensile at 5%, g/in	1336	1597	1370	1659
MD Tensile at 10%, g/in	2837	3691	2758	3686
MD Tensile at 25%, g/in	5598	—	4736	5025
MD Tensile at Break, g/in	9294	9934	6131	7479
MD Elong. at Break, %	78.35	21.08	75.56	24.07
TD Tensile at 5%, g/in	303.9	121.3	241.8	144.2
TD Tensile at 10%, g/in	473.4	238.2	379.2	245.6
TD Tensile at 25%, g/in	589.7	421.6	473.8	326.9
TD Tensile at Break, g/in	820.8	464.8	634.7	356.5
TD Elong. at Break, %	388.0	330.2	356.8	270.3
MD Elmendorf Tear, grams	0	0	13.2	13.2
MD Shrink at 170° F, %	13	18	11.5	14.9
TD Shrink at 170° F, %	-3	-3	-3	-2.5

Note: All samples oriented with a 15 fpm inlet speed, 190° F annealing and 5% relaxation.

**TABLE 6**

PROPERTY		Example 24 mLLDPE 50% CaCO <sub>3</sub> 2.7:1 draw ratio	Example 25 Z-N LLDPE 50% CaCO <sub>3</sub> 3.8:1 draw ratio
Yield	yd <sup>2</sup> /lb	23.62	26.23
Basis Weight	g/m <sup>2</sup>	23.13	20.85
Embossed Caliper	mils	1.26	1.61
Gurley Porosity	Seconds/100 cc	251	230
WVTR	g/m <sup>2</sup> /24 hours	7613	7688
MD Tensile at 5% Elg.	grams/in	195.5	174.7
MD Tensile at 10% Elg.	grams/in	269.1	272.9
MD Tensile at 25% Elg.	grams/in	301.7	321.8
MD Tensile at Break	grams/in	477.6	431.7
MD Elong. at Break	%	346.4	293.7
TD Tensile at 5% Elg.	grams/in	371.5	553.3
TD Tensile at 10% Elg.	grams/in	622.0	980.4
TD Tensile at 25% Elg.	grams/in	932.9	1702
TD Tensile at Break	grams/in	1650	2162
TD Elong. at Break	%	116.5	86.4
TD Shrinkage at 170°F	%	4.2	4.0

**CLAIMS**

We Claim:

1. A process of making a housewrap, comprising:
  - 5 a) mixing a polyolefin with a filler;
  - b) extruding a film from the polyolefin/filler mixture;
  - c) melt embossing the precursor film to impose therein a pattern of different film thickness
  - d) stretching the film of c) to impart greater WVTR; and
  - 10 e) laminating the stretched film to a nonwoven fabric comprising cross-laminated fibers at a temperature and pressure sufficient to bond the fabric and film to form a breathable laminate, characterized that said polyolefin includes at least a majority of a polyethylene having a  $M_w/M_n < 3$ , and a CDBI above 60 percent;
- 15 wherein the film will have a WVTR at least 10% higher than the WVTR described by the equation: 
$$\text{WVTR} = -10,900 + 320 (\text{filler weight } \%)$$
2. The method of claim 1, further comprising heat setting the stretched film at a temperature above the stretching temperature and below the softening  
20 temperature of the stretched film;
  - wherein said lamination is selected from the group consisting of heat lamination, adhesive lamination, extrusion lamination, mechanical bonding and combinations thereof;
  - wherein the polyolefin is a copolymer of ethylene and a  $C_4$ - $C_{10}$  alpha-olefin  
25 having a  $M_w/M_n < 2.5$ ;
  - wherein said filler is calcium carbonate surface treated with calcium stearate;
  - wherein the precursor film is melt embossed with a diamond pattern; and
  - wherein the polyolefin/filler mixture contains between 30 percent to 65  
30 percent filler by weight based on the total weight of said mixture.

3. The method of any of the preceeding claims, wherein the fabric is a nonwoven polyolefin fabric having a heat seal layer.
- 5 4. The method of claim 3, wherein the lamination comprises heating the fabric and pressing the unheated film to the heated fabric.
5. The process of any of the previous claims, wherein the polyolefin/filler mixture contains between 30 percent to 65 percent filler by weight.
- 10 6. The process of any of the preceeding claims, wherein said film has a WVTR of at least 20% greater than the WVTR described by the equation:
- $$\text{WVTR} = -10,900 + 320 (\text{filler weight } \%)$$
- 15 7. The process of any of the preceeding claims, wherein said film has a WVTR of at least 30% greater than the WVTR described by the equation:
- $$\text{WVTR} = -10,900 + 320 (\text{filler weight } \%)$$

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## AMENDED CLAIMS

[received by the International Bureau on 20 January 1998 (20.01.98);  
new claim 8 added; remaining claims unchanged (1 page)]

3. The method of any of the preceeding claims, wherein the fabric is a nonwoven polyolefin fabric having a heat seal layer.
- 5 4. The method of claim 3, wherein the lamination comprises heating the fabric and pressing the unheated film to the heated fabric.
5. The process of any of the previous claims, wherein the polyolefin/filler mixture contains between 30 percent to 65 percent filler by weight.
- 10 6. The process of any of the preceeding claims, wherein said film has a WVTR of at least 20% greater than the WVTR described by the equation:  
$$\text{WVTR} = -10,900 + 320 (\text{filler weight } \%)$$
- 15 7. The process of any of the preceeding claims, wherein said film has a WVTR of at least 30% greater than the WVTR described by the equation:  
$$\text{WVTR} = -10,900 + 320 (\text{filler weight } \%)$$
8. A housewrap made from any of the preceeding claims.

1 / 1

FIG. 1

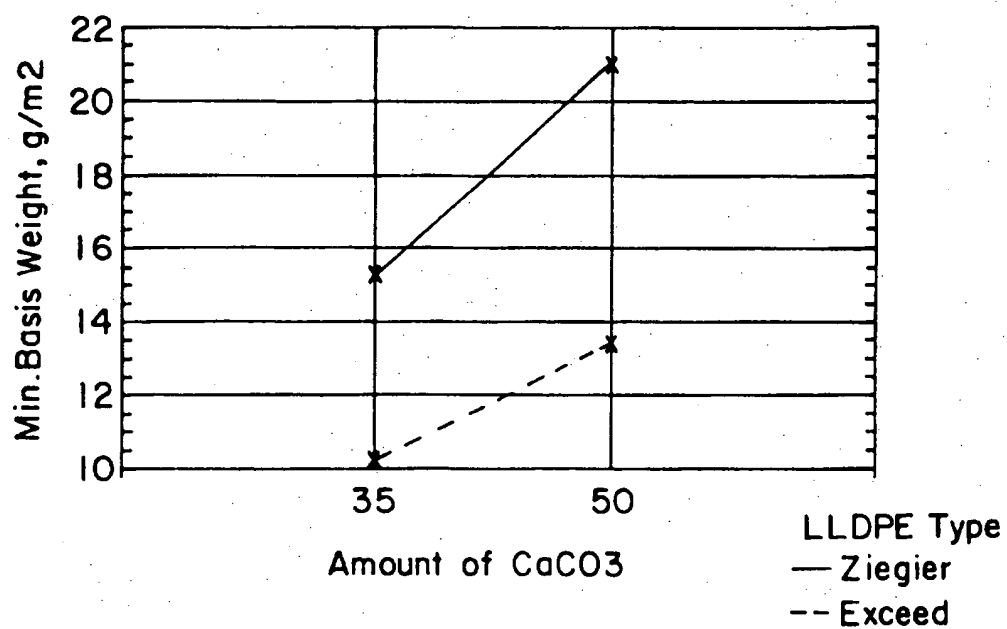
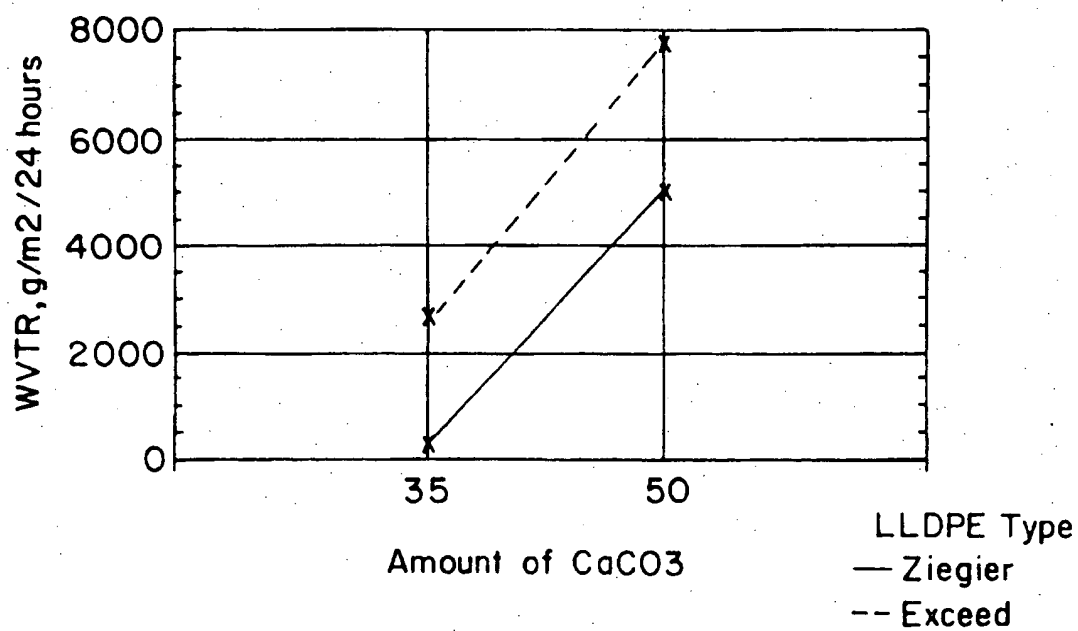


FIG. 2



# INTERNATIONAL SEARCH REPORT

Intern Application No

PCT/US 97/13580

## A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 B32B31/12 B32B27/12

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 B32B

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

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Y	US 4 929 303 A (SHETH PARESH J) 29 May 1990 see column 3, line 7-26 - column 3, line 59-63; claims 1-17 ---	1-7
Y	PATENT ABSTRACTS OF JAPAN vol. 095, no. 008, 29 September 1995 & JP 07 118431 A (MITSUBISHI CHEM CORP), 9 May 1995, see abstract ---	1-7
Y	EP 0 691 203 A (KIMBERLY CLARK CO) 10 January 1996 see claims 1,9,10; example 1 ---	1-7
A	US 5 169 712 A (TAPP WILLIAM T) 8 December 1992 see claims 1,6,15-18 ---	1-7
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☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

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Date of the actual completion of the international search

11 November 1997

Date of mailing of the international search report

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# INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 97/13580

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
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